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(21) International Application Number: PCT/US93/02431 (22) International Filing Date: 18 March 1993 (18.03.93) (30) Priority data: 854,125 19 March 1992 (19.03.92) US (71) Applicant: ENGELHARD CORPORATION [US/US]; 101 Wood Avenue, Iselin, NJ 08830 (US). (72) Inventors: MADON, Rostam, J. ; 9 Hendrick Road, Flemington, NJ 08822 (US). KOERMER, Gerald, S. ; 37 Stonegate Drive, Roseland, NJ 07068 (US). DEEBA, Michael ; 2501 North Oaks Boulevard, North Brunswick, NJ 08902 (US). YAU, Kathleen, Y. ; 3 Van Horn Place, Holland, PA 18966 (US).	(74) Agents: MILLER, Stephen, I. et al.; Engelhard Corporation, 101 Wood Avenue, Iselin, NJ 08830 (US). (81) Designated States: CA, JP, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published With international search report.	
(54) Title: FLUID CATALYTIC CRACKING PROCESS FOR INCREASED FORMATION OF ISOBUTYLENE AND ISOAMYLENES (57) Abstract A composite fluid catalytic cracking catalyst comprising a non-zeolitic component of at least 45 % Al ₂ O ₃ and containing ≤ 30 % Y zeolite with unit cell size ≤ 24.29 Å, the matrix (non-zeolitic component) of said catalyst having a B.E.T. surface area ≥ 50 m ² /g and a catalyst Lewis/Bronsted acid site ratio of ≥ 1.5 wherein the total number of Bronsted acid sites is ≤ 30 micromoles/g of catalyst. The catalyst is used to increase the isobutylene and isoamylenes content of cracked products obtained in a fluid catalytic cracking unit.		

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FLUID CATALYTIC CRACKING PROCESS FOR INCREASED
FORMATION OF ISOBUTYLENE AND ISOAMYLENES

BACKGROUND OF THE INVENTION

5 This invention relates to the fluid catalytic cracking (FCC) of petroleum feedstocks to produce transportation fuels and is especially directed to improvements in FCC to increase the isobutylene and isoamylenes contents of the cracked product.

10 Increased addition of oxygenated materials to gasoline has been mandated by the US Government. The current industry additives of choice are methyl tertiary butyl ether (MTBE) and tertiary amyl methyl ether (TAME). The technology to make MTBE is well known and
15 commercially viable. Briefly, the ether is manufactured by reacting isobutylene and methanol over a resin catalyst. Obtaining sufficient amounts of isobutylene and isoamylenes at reasonable cost is crucial. In most modern refineries, a dehydroisomerization process can be
20 operated to convert butanes to isobutylene. However, a fluid catalytic cracker is the best if not the only source of making isobutylene and isoamylenes while processing heavy hydrocarbon streams such as gas oil and resid. An FCC catalyst that enhances these olefin
25 yields is essential for viable production of MTBE and TAME in refineries.

 It is well known in the literature that lowering the unit cell size (UCS) of a Y zeolite via steaming gives a catalyst which enhances olefin selectivity
30 during the catalytic cracking of hydrocarbons. We have discovered, however, that we can make further improvements in isobutylene and isoamylenes yields than by just lowering unit cell size of Y.

SUMMARY OF THE INVENTION

The present invention comprises an FCC operation wherein the catalyst is a composite comprising a non-zeolitic alumino-silicate (or mixtures of Al_2O_3 , SiO_2 , and $\text{SiO}_2\text{-Al}_2\text{O}_3$ component) of at least 45% Al_2O_3 and containing $\leq 30\%$ Y zeolite with a unit cell size $\leq 24.29 \text{ \AA}$, the matrix of said catalyst having a BET surface area $> 50 \text{ m}^2/\text{g}$, said composite catalyst having a Lewis/Bronsted acid site ratio of > 1.5 , and preferably greater than 2.5, wherein the total number of Bronsted acid sites is < 30 micromoles/g of catalyst, preferably less than 20 micromoles/g of catalyst, but greater than 0 micromoles/g of catalyst.

Such composite catalysts give significantly enhanced amounts of isobutylene and isoamylenes during catalytic cracking of gas oil. The % Y zeolite is determined by x-ray diffraction (ASTM D3906-85a). Unit cell size is determined by x-ray diffraction using ASTM D3942-85.

THE DRAWINGS

Figure 1 is a plot of relative isobutylene yield vs. activity [conversion/(100-conversion)] for a standard commercial FCC catalyst and a catalyst used in practice of the invention.

Figure 2 is a plot of effect of time of hydrothermal treatment in a preparation of a catalyst used in the invention.

DESCRIPTION OF PREFERRED EMBODIMENTS

The starting material used to prepare catalysts employed in practice of this invention may be an in situ type FCC catalyst supplied by Engelhard Corporation such as MONADYNE™, PRECISION™ or OCTIDYNE™ catalysts. The starting material may also be a Y zeolite which has been intimately bound with an amorphous alumino-silicate matrix. It is important that the amorphous part of the composition is high in aluminum content which should be greater than 45 wt%. The Al_2O_3 may be part of an alumino-silicate material or mixed with silica, or mixed with silica-alumina. The Al_2O_3 mixed with the above amorphous components may be any phase of Al_2O_3 . It is preferable to have minimal (e.g., less than 1.0% ReO by weight) and preferably no rare earth cations and to have sodium levels below 0.3 wt% Na_2O , preferably 0.1% or below, based on the dry weight of catalyst.

Catalysts of the invention may be made by severe hydrothermal treatment of the types of starting materials stated above. Hydrothermal treatment conditions will depend on the amount of Y zeolite in the starting material. In general, the conditions include (i) temperatures of 1450°F to 1800°F (ii) times of 1 to 25 h, and (iii) 25 to 100% steam. Higher temperatures may require shorter times and lower steam partial pressure. For example, at 1450°F, Material A (described below) requires 15 to 24 h of hydrothermal treatment in 100% steam; whereas at 1550°F only approximately 5 hours are sufficient to get similar results. Hydrothermal treatment does not affect the amount of rare earth or sodium cations which are low as stated above for the starting materials.

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Additive materials compatible with FCC processing such as octane boosters (ZSM-5) and Beta, CO combustion promoters, SOx capture additives, etc., may be added to such catalysts.

5 The FCC process of the invention may be carried out in conventional heat balanced FCC units in the absence of added hydrogen under conditions which will vary depending upon the refinery. Operating variables for the riser and regenerator are well known in the art and
10 are described extensively in the literature. See, for example, CATALYSIS REVIEWS - SCIENCE AND ENGINEERING, VOL. 18, NUMBER 1, 1978. While conventional FCC cracking and regeneration conditions may be used with
15 conventional FCC feedstock such as gas oils and resids, catalysts of the invention produce a product slate that is different from those achieved with conventional FCC catalysts. Most significant are the appreciable increases in isobutylene and isoamylene yields. Gasoline yields decrease whereas octane number
20 increases. Isobutylene and isoamylenes are recovered from cracked products using conventional FCC separation technology.

 We have also found that addition of relatively large amounts (e.g., 10-20% by weight of kaolin bound
25 microspheres containing about 15% ZSM-5) to our isobutylene catalysts results in an unexpected significant decrease in coke yield and an increase in activity.

 The performance properties of modified catalysts of
30 the invention are such that some changes in FCCU operation may be required in order to take full advantage of the potential yield benefits of the catalysts. In particular, the total wet gas (C₃ and C₄ hydrocarbons) handling capacity of the unit may need to
35 be increased. This can be accomplished by means well known to those skilled in the art.

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Commercial yield projections for various operating scenarios for maximizing light olefin yields were carried out by computer simulation. These projections indicated that isobutylene yield increases of 50-200% can be achieved compared with a typical current operation. The results show that catalysts of the invention are more selective for increasing isobutylene yield than other options such as high riser temperatures or ZSM-5 additives with conventional FCC catalysts.

While it is expected that some investment in the gas plant of an FCCU will be required, this may turn out to be an economically preferred route compared with alternative supplies of isobutylene produced by isomerization and dehydrogenation of butanes.

In the following illustrative examples, the starting cracking catalysts material described as Material A contained approximately 20 wt% Y, 0% rare earth, and < 0.1% Na₂O; material B contained approximately 40 wt% Y, 0.1% rare earth, and 0.1% Na₂O; and material C contained approximately 12 wt% Y, 0% rare earth, and < 0.3% Na₂O. In all these materials, the rest of the composition, and thus the predominant part of the starting material is amorphous alumino-silicate, in which the total Al₂O₃ content was greater than 50%.

Above materials when hydrothermally treated are designated AH, BH, and CH, respectively. Hydrothermal treatments of materials in the examples were performed with 100% steam at ambient pressure.

Catalyst testing and hydrothermal treatments were carried out using procedures referred to in U.S. 4,493,902, Brown et al unless otherwise indicated. All yields given in the examples are on a weight-on-feed-basis.

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Lewis and Bronsted acidity values were measured as follows. Infrared spectra were collected at 40°C in the diffuse reflectance mode using a Spectra Tech controlled-environment chamber in a Perkin Elmer 1750 spectrometer. The samples were dried for 1h at 723 K under flowing N₂, and the dehydration process was monitored by observing the disappearance of the water band at 1645 cm⁻¹. Pyridine vapor was then introduced to the sample in flowing N₂ at room temperature; after 1 min the flow was shut off and the sample allowed to equilibrate with the pyridine for 1h. Spectra collected after each of two successive 1h purges in N₂ at 40 and 200°C indicated two distinct desorption stages. The lowest temperature purge removed predominantly physisorbed pyridine (1583 cm⁻¹) while the second eliminated hydrogen-bonded pyridine (1600 cm⁻¹). Pyridine was not decomposed at the highest desorption temperature. Infrared band areas (Kubelka-Munk units) at 1450 and 1550 cm⁻¹ were used to estimate the amounts of pyridine adsorbed on Lewis and Bronsted acid sites, respectively. These areas were reproducible to within about 10% due to variations in sample grinding and packing in the reflectance chamber sample cup.

Unlike conventional quantitative transmission infrared spectroscopy, where extinction coefficients are obtained from a linear plot of band intensity versus sample concentration, there is no clearly defined equivalent "beam scatter" parameter for the diffuse reflectance technique. This makes defining an

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extinction coefficient in diffuse reflectance subjective; it includes contributions from the experimental setup and the sample density which are difficult to quantify. In this study, extinction

5 coefficients have been estimated by correcting errors due to sample density differences by using both ammonia desorption and thermogravimetric pyridine adsorption techniques to obtain total acidity estimates for our samples. Therefore the extinction coefficients used to

10 quantify our data were effective extinction coefficients, and these values may not be directly transferable to a different experimental setup. Their relative values, however, will be constant, allowing quantitative comparison between the various samples of

15 this study.

Acidity, in equivalent mmol/g, was calibrated using ammonia capacity at 200°C to determine total acidity for Lewis only (alumina) and Bronsted-only (pure H-zeolites) samples. This calibration converts 200°C infrared

20 pyridine peak areas in units of Kubelka Munk reflectance to equivalent ammonia acidity. The Pyridine peak for Bronsted acid sites at 1550 cm^{-1} divided by 4.7 gives equivalent Bronsted acidity. The pyridine peak for Lewis acid sites at 1450 cm^{-1} divided by 6.9 gives

25 equivalent Lewis acidity.

EXAMPLE 1

The following table gives results of microactivity tests (MAT) at 910°F and 48s gas oil delivery time.

Catalyst/oil ratios were adjusted to obtain 70%

5 conversion.

TABLE 1

	<u>MATERIAL</u> <u>AH</u>	<u>MATERIAL</u> <u>BH</u>	<u>MATERIAL</u> <u>B</u>
10 HYDROTHERMAL TREATMENT CONDITIONS	1550°F/5h	1550°F/5h	1450°F/4h
UCS, A	24.26	24.27	24.29
CATALYST/OIL	8.1	6.3	5
CONVERSION, %	71	70	71
15 C ₄ YIELDS, %			
Isobutylene	1.8	1.6	1.0
Straight butenes	3.8	3.7	2.9
Isobutane	4.5	4.5	5.5
N-butane	0.7	0.8	0.9
20 <u>Isobutylene</u>			
Total C ₄ olefins	0.32	0.30	0.26
<u>C₄ Olefins</u>			
C ₄ paraffins	1.1	1.0	0.6

Data in Table 1 show that Materials AH and BH,
 25 hydrothermally treated at the more extreme conditions,
 gave higher yields of isobutylene and olefins than
 Material B steamed at conventional conditions (1450°F,
 4h).

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Table 2 compares Material AH with a commercial HY zeolite, both hydrothermally treated to the same extent. The data show that even though Material AH has a higher unit cell size than steamed commercial Y, it made significantly more isobutylene than the commercial Y. Also, for Material AH isobutylene was closer to the thermodynamic equilibrium ratio of 0.45.

TABLE 2

	<u>MATERIAL AH</u>	<u>COMMERCIAL Y ZEOLITE</u>
10	1500°F/24h	1500°F/25h
	HYDROTHERMAL CONDITIONS	
	UCS, A	24.27
	CATALYST/OIL RATIO	24.23
	CONVERSION, %	5
		66
15	C ₄ YIELDS, %	
	Isobutylene	1.9
	Straight butenes	0.6
	Isobutane	3.7
	N-butane	4.1
		4.2
		0.7
20	<u>Isobutylene</u>	
	Total C ₄ olefins	0.34
		0.27
	<u>C₄ Olefins</u>	
	C ₄ paraffins	1.2
		0.45

EXAMPLE 2

Figure 2 shows the effect of time of hydrothermal treatment on Material C. Catalyst/oil ratios were varied to give 72% MAT conversions for all hydrothermally treated catalysts. Note that after 4 h steaming at 1450°F, the UCS was already 24.27 A, with an isobutylene yield of 1.3%; further steaming up to 60 h increased the isobutylene yield significantly to 2.8 % even though the UCS only changed to 24.26 A.

10 EXAMPLE 3

Figure 1 is a plot of relative isobutylene yield vs. activity [conversion/(100-conversion)] for two FCC catalysts. One was a standard commercial FCC catalyst. The other was a typical catalyst of this invention. The data was generated from the standard microactivity test where activity was varied by changing the catalyst to oil ratio and the wt% isobutylene yield on feed was divided by a constant. The standard commercial catalyst gave a very small increase in isobutene as activity increased. In contrast, the isobutene yield of the catalyst of the present invention increased dramatically as activity increased.

EXAMPLE 4

Material A was hydrothermally treated for 7h at 1550°F using 100% steam at 1 bar to give AH. This material was divided into two portions. One portion was run in the microactivity test. The second portion was blended with ZSM-5 in a kaolin matrix so that the % ZSM-5 in the composite was about 15% and the ZSM-5 in the total catalyst was about 3%. This blend was also tested by the standard microactivity test.

10 The results are in Table 3. Data in this table show that the coke yield from the catalyst containing ZSM-5 was lower than that of the catalyst without ZSM-5. In addition the activity of the catalyst with ZSM-5 was higher than the catalysts without ZSM-5 at the same catalyst to oil ratio. With ordinary cracking catalyst the typical behavior with regard to activity is that activity decreases by approximately the blend ratio of the additive; i.e., a catalyst containing 20% of the ZSM-5 blend would have almost a 20% decrease in activity. This normally occurs because ZSM-5 has a very low primary cracking activity.

EXAMPLE 5

Material A was hydrothermally treated at 1500°F for 10h with 100% steam at 1 bar to obtain AH. This catalyst was tested by running a standard MAT. In addition, the hydrothermally treated catalyst was blended with enough ZSM-5 blend to give a blend containing 80% AH. This catalyst blend was also tested by a standard MAT. The catalyst without ZSM-5 had higher coke yields and lower activity. See Table 4.

10. EXAMPLE 6

In the following example, pilot unit studies were carried out with the same gas oil feed as used in the MAT on a modified recirculating ARCO LAB - FCC pilot unit (W.H. Humes, Chemical Eng. Progress, page 51, Feb., 1983) which has a 17 foot lean phase plug flow cracking riser. The unit was operated under computer control at atmospheric pressure and nitrogen was used for dispersion. The catalyst charge to the unit was around 2600 g with an oil feed rate between 9 and 12 g/min.

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The riser was operated under adiabatic conditions with the riser temperature at the top being 960°F. Catalyst regeneration was carried out in the regenerator at 1200°F with air at 9 liters/min. Conversion was adjusted by
5 varying catalyst circulation rate and/or oil feed rate. Liquid products and unreacted gas oil were separated from product gases with the help of a C₄-C₅ fractionation column. The product gases and flue gases were analyzed by standard gas chromatography. The liquid was analyzed
10 by simulated distillation chromatography (ASTM D2887). A more detailed breakdown of the gasoline fraction was obtained using a PIONA analyzer from Analytical Controls. This latter analysis allowed us to obtain the concentration of isoamylenes in the gasoline.

15 A catalyst composite was made as follows. Material A was hydrothermally treated at 1500°F for 10 h. Material B was hydrothermally treated at 1450°F for 23 h. The hydrothermally treated materials were blended as follows: 87.5 wt% AH and 12.5 wt% BH. This resulting
20 catalyst was run in the recirculating FCC pilot unit. At 68% conversion, the gasoline fraction contained 6.89 wt% isoamylenes.

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TABLE 3

Effects of ZSM-5 on High Isobutylene Catalysts

Catalyst AH

	C/O	5.03	4.96	7.10	7.07
5	Activity	1.22	1.42	1.56	1.59
	Coke Yield	4.65	4.59	5.18	5.13
	Coke/Act	3.81	3.23	3.32	3.23

Catalyst AH + 16% ZSM-5 Blend

	C/O	5.06	5.00	6.99	7.08	7.04	7.07
10	Activity	1.53	1.54	1.74	1.76	1.74	1.82
	Coke Yield	4.04	4.04	4.76	4.77	4.76	4.91
	Coke/Act	2.64	2.62	2.74	2.71	2.74	2.70

TABLE 4

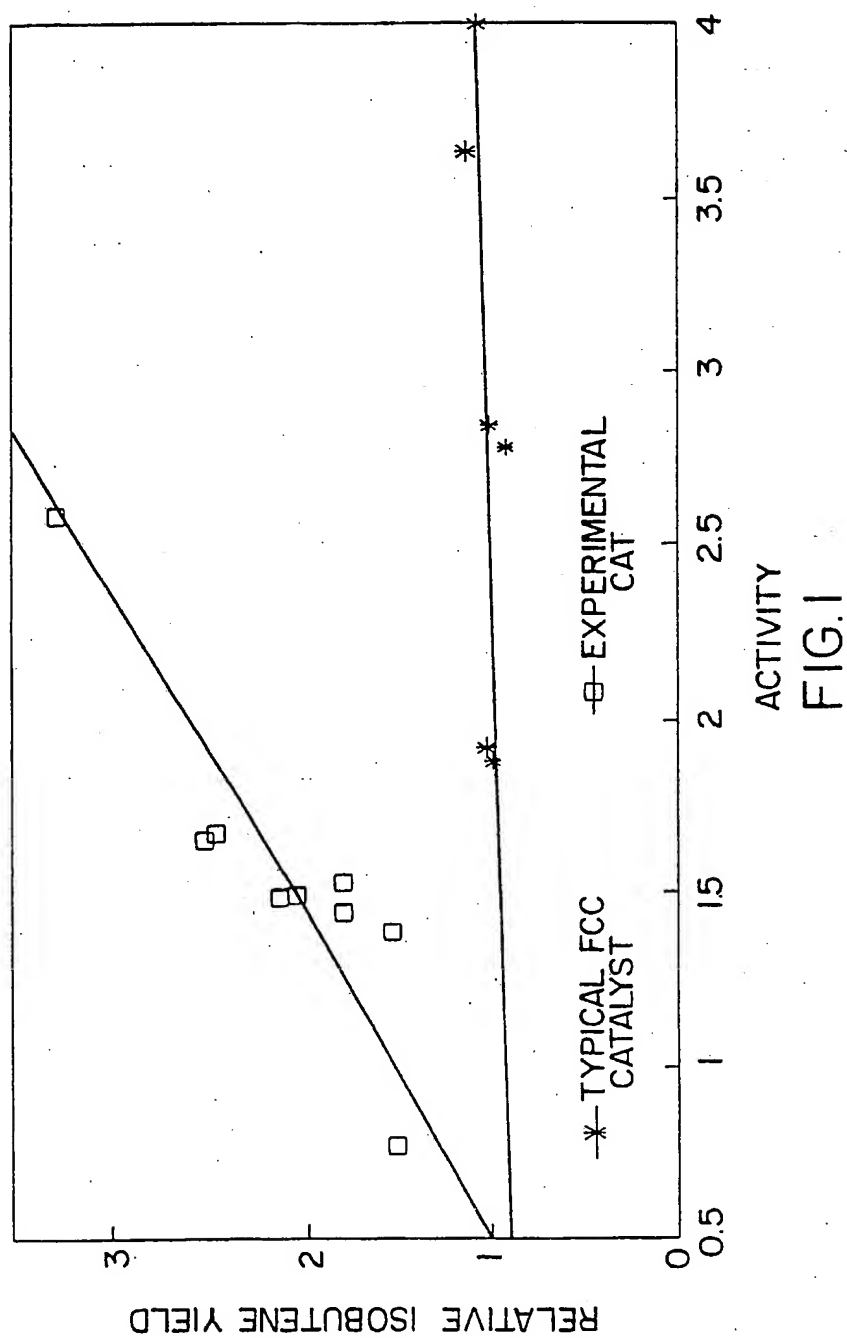
Effects of ZSM-5 on Coke Yield

Catalyst		<u>AH</u>			<u>AH + 20% ZSM-5 Blend</u>		
5	C/O	4.99	5.04	5.01	5.08	7.13	7.04
	Activity	1.23	1.66	1.41	1.59	1.95	2.06
	Coke	4.00	5.17	3.89	3.94	4.96	4.86
	Coke/Activity	3.25	3.11	2.76	2.48	2.54	2.36

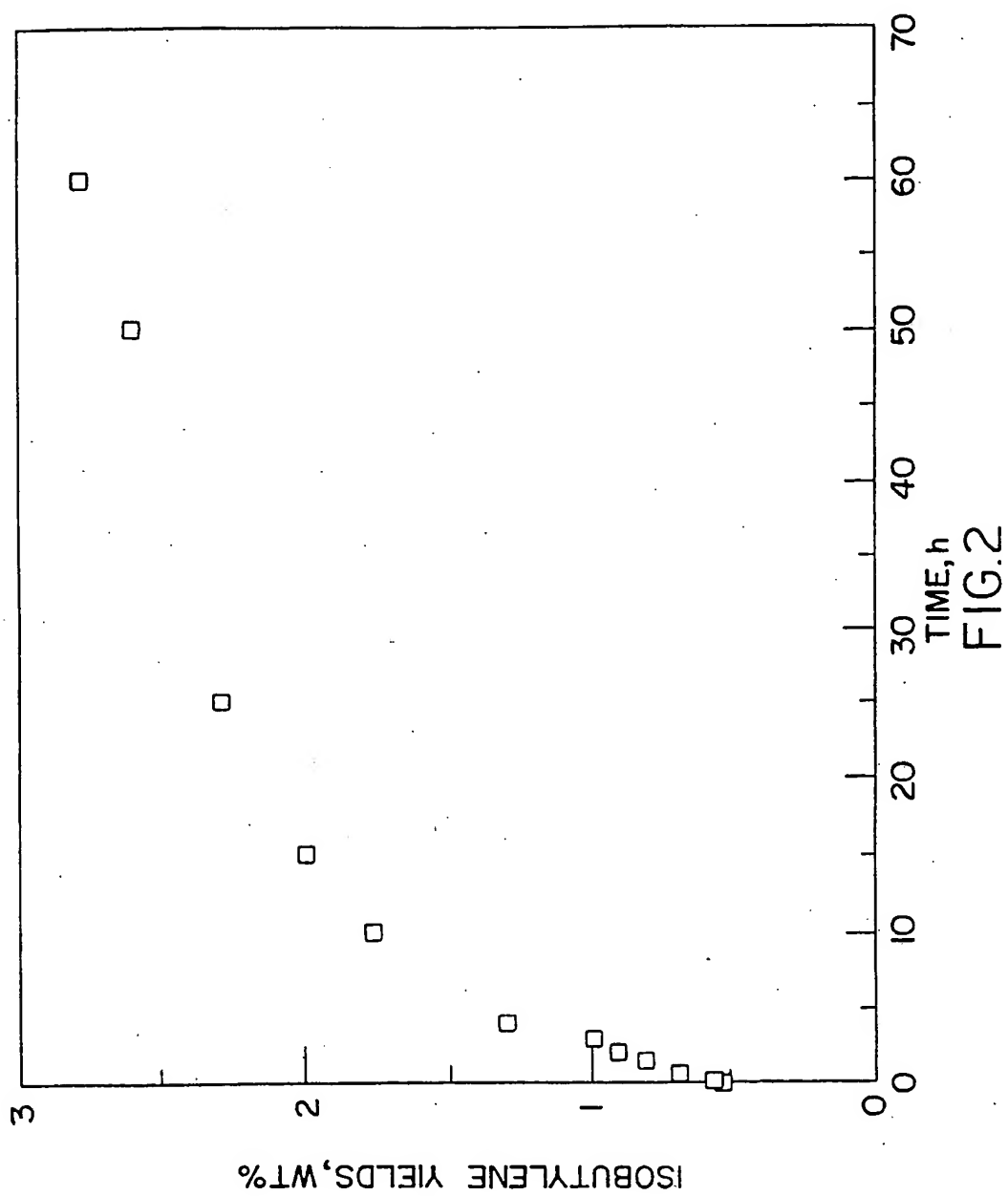
We claim:

1. A process for the catalytic cracking of a hydrocarbon charge to produce isobutylene and isoamylenes which comprises contacting said hydrocarbon charge at cracking temperature in the absence of added hydrogen with a composite fluid catalytic cracking catalyst comprising a non-zeolitic component containing at least 45% by weight Al_2O_3 and $\leq 30\%$ Y zeolite having a unit cell size $\leq 24.29 \text{ \AA}$, the matrix of said composite catalyst having a BET surface area $\geq 50 \text{ m}^2/\text{g}$, said catalyst having a Lewis/Bronsted acid site ratio of ≥ 1.5 wherein the total number of Bronsted sites is ≤ 30 micromoles/g of catalyst, and recovering from the resulting products of conversion a concentrate of isobutylenes and isoamylenes.
2. The process of claim 1 wherein the total number of Bronsted sites is less than 20 micromoles/g of catalyst.
3. The process of claim 1 wherein said catalyst contains less than 0.3% Na_2O on a dry weight basis.
4. The process of claim 1 wherein said catalyst contains less than 0.1% Na_2O on a dry weight basis.
5. The process of claim 1 wherein said catalyst contains less than 1% REO on a dry weight basis.
6. The process of claim 1 wherein said catalyst is essentially free from rare earth.
7. The process of claim 1 wherein the Al_2O_3 content of said non-zeolitic component is selected from the group consisting of alumino-silicate, silica-alumina, mixtures of alumina and silica, and mixtures thereof.
8. The process of claim 1 wherein ZSM-5 is present with said cracking catalyst.
9. The process of claim 1 wherein the unit cell size of said Y zeolite catalyst is less than 24.26 Angstroms.

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INTERNATIONAL SEARCH REPORT

PCT/US 93/02431

International Application No

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) ⁶		
According to International Patent Classification (IPC) or to both National Classification and IPC		
Int.Cl. 5 C10G11/05		
II. FIELDS SEARCHED		
Minimum Documentation Searched ⁷		
Classification System	Classification Symbols	
Int.Cl. 5	C10G ; C07C	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched ⁸		
III. DOCUMENTS CONSIDERED TO BE RELEVANT⁹		
Category ¹⁰	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³
A	WO,A,8 603 139 (UNION CARBIDE) 5 June 1986 see claims 1,3,7,13,23 see page 6, line 20 - line 25 see page 7, line 16 - page 40, line 10 see page 41, line 1 - line 18 see page 43, line 19 - line 22 see page 47, line 8 - line 18 ----	1,7,8
A	US,A,5 001 097 (CHEVRON) 19 March 1991 see claims 1,3,5,6,7,22 see column 7, line 52 - line 59 ----	1,3,7
A	EP,A,0 256 875 (MOBIL) 24 February 1988 -----	
<p>¹⁰ Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"A" document member of the same patent family</p>		
IV. CERTIFICATION		
Date of the Actual Completion of the International Search	Date of Mailing of this International Search Report	
07 JUNE 1993	18. 06. 93	
International Searching Authority	Signature of Authorized Officer	
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US 9302431
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